ABSTRACT

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A method and composition for the prevention, amelioration or control of external parasites on animals and humans utilizing a compound of formula I.

$$(R)_{n}$$

$$R_{1}$$

$$R_{2}$$

$$R_{3}$$

$$R_{4}$$

$$(I)$$

5 or a pharmaceutically acceptable salt thereof, wherein

R is halogen, OR₇, SO_mR₈, NO₂, CN, C₁-C₆haloalkyl or an optionally substituted C₁-C₆alkyl group;

n is 0 or an integer of 1, 2 or 3;

m is 0 or an integer of 1 or 2;

R₁ is H, halogen, NO₂, NR₉R₁₀, NR₁₁COR₁₂, NCHNR₉R₁₀ or NCHOR₁₃;

R₂, R₃, R₄, R₅ and R₆ are each independently H, halogen or a C₁-C₄alkyl, aryl or heteroaryl group each optionally substituted;

R₇ is H or a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, aryl or heteroaryl group each optionally substituted;

R₈ is a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, aryl or heteroaryl group, each optionally substituted;

 R_9 and R_{10} are each independently H, C_1 - C_4 haloalkyl or a C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, aryl or heteroaryl group each optionally substituted or R_9 and R_{10} may be taken together with the atom to which they are attached to form a 5- to 7-membered ring optionally containing 1 or 2 additional heteroatoms selected from O, N or S;

R₁₁ is H, COR₁₂ or an optionally substituted C₁-C₄alkyl group;

R₁₂ is a C₁-C₆alkyl, C₂-C₆alkenyl, C₂-C₆alkynyl, aryl or heteroaryl group each optionally substituted; and

R₁₃ is H or a C₁-C₆alkyl, aryl or heteroaryl group each optionally substituted; or a stereoisomer or tautomer thereof.